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HOUSE BILL 479

47TH LEGISLATURE - STATE OF NEW MEXICO - FIRST SESSION, 2005

INTRODUCED BY

Rhonda S. King

AN ACT

RELATING TO HEALTH; EXPANDING THE NUMBER OF MEDICAL TESTS  
REQUIRED FOR NEWBORN CHILDREN.

BE IT ENACTED BY THE LEGISLATURE OF THE STATE OF NEW MEXICO:

Section 1. Section 24-1-6 NMSA 1978 (being Laws 1973,  
Chapter 359, Section 6, as amended) is amended to read:

"24-1-6. TESTS REQUIRED FOR NEWBORN INFANTS. --

A. The department shall adopt tests for the  
detection of [~~phenylketonuria and other~~] congenital diseases  
[~~which~~] that shall be given to every newborn infant, except  
that, after being informed of the reasons for the tests, the  
parents or guardians of the newborn child may waive the  
requirements for the tests in writing. The tests shall include  
at a minimum:

- (1) 3-methylcrotonyl-CoA deficiency;

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- 1                                    (2) 3-OH 3-CH<sub>3</sub> glutaric aciduria;
- 2                                    (3) argininosuccinic acidemia;
- 3                                    (4) beta-ketothiolase deficiency;
- 4                                    (5) biotinidase deficiency;
- 5                                    (6) carnitine uptake defect;
- 6                                    (7) citrullinemia;
- 7                                    (8) congenital adrenal hyperplasia;
- 8                                    (9) congenital hypothyroidism;
- 9                                    (10) cystic fibrosis;
- 10                                   (11) galactosemia;
- 11                                   (12) glucose-6-phosphate dehydrogenase
- 12                                   deficiency;
- 13                                   (13) glutaric acidemia type I;
- 14                                   (14) Hb S/beta-thalassemia;
- 15                                   (15) Hb S/C disease;
- 16                                   (16) hearing deficiency;
- 17                                   (17) homocystinuria;
- 18                                   (18) isovaleric acidemia;
- 19                                   (19) long-chain L-3-OH acyl-CoA dehydrogenase
- 20                                   deficiency;
- 21                                   (20) maple syrup urine disease;
- 22                                   (21) medium chain acyl-CoA dehydrogenase
- 23                                   deficiency;
- 24                                   (22) methylmalonic acidemia;
- 25                                   (23) methylmalonic acidemia (mutase

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1 deficiency);

2 (24) multiple carboxylase deficiency;

3 (25) phenylketonuria;

4 (26) propionic acidemia;

5 (27) sickle cell anemia;

6 (28) trifunctional protein deficiency;

7 (29) tyrosinemia type I; and

8 (30) very long-chain acyl-CoA dehydrogenase

9 deficiency.

10 B. In determining which other congenital diseases  
11 to adopt tests for, the secretary [~~of health and environment~~]  
12 shall consider the recommendations of the New Mexico pediatrics  
13 society of the American academy of pediatrics.

14 [~~B.~~] C. The department shall institute and carry on  
15 such laboratory services or may contract with another agency or  
16 [~~state~~] entity to provide such services as are necessary to  
17 detect the presence of [~~phenylketonuria and other~~] congenital  
18 diseases.

19 [~~C.~~] D. The department shall, as necessary, carry  
20 on an educational program among physicians, hospitals, public  
21 health nurses and the public concerning [~~phenylketonuria and~~  
22 ~~other~~] congenital diseases.

23 [~~D.~~] E. The department shall require that all  
24 hospitals or institutions having facilities for childbirth  
25 perform or have performed tests for [~~phenylketonuria and other~~]

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underscored material = new  
~~[bracketed material] = delete~~

1 congenital diseases on all newborn infants except if the  
2 parents or guardians of a child object to the tests in  
3 writing. "

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